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MEMORANDUM FOR PRS (In-House Publication)

FROM: PROI (STINFO)

10 April 2001

SUBJECT: Authorization for Release of Technical Information, Control Number: AFRL-PR-ED-VG-2001-078
Fajardo, Mario, "Chemistry and Spectroscopy in Solid Parahydrogen"

U. Wyoming Chemistry Dept. Seminar
(Caramie, WY, 20 April 2001) (Deadline: 20 April 2001)

(Statement A)

1. This request has been reviewed by the Foreign Disclosure Office for: a.) appropriateness of distribution statement, b.) military/national critical technology, c.) export controls or distribution restrictions, d.) appropriateness for release to a foreign nation, and e.) technical sensitivity and/or economic sensitivity.

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APPROVED/APPROVED AS AMENDED/DISAPPROVED

PHILIP A. KESSEL
Technical Advisor
Space & Missile Propulsion Division

Date

Chemistry and Spectroscopy in Solid Parahydrogen

Mario E. Fajardo

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- * Cryosolid Propellants Team
- * HEDM Cryosolid Propellants Concept (Atoms in Solid Hydrogen)
- * Rapid Vapor Deposition of Transparent Parahydrogen (pH₂) Solids
- * B and Al Doped pH₂ Solids
- * High Res. IR Spectroscopy of Molecular Dopeants in Solid pH₂
- * Summary

Cryosolid Propellants Team

* Mario E. Fajardo, Michelle E. DeRose, and Simon Tam

* Bill Larson (thermal B atom source)

* Jeff Sheehy, Jerry Boatz, Peter Langhoff (in-house theory)

* AFOSR Contractors:

- P. Dagdigian @ Johns Hopkins: Al/H₂ & B/H₂ Complexes
- M. Alexander @ U. Maryland: B/H₂ Interaction Potentials
- G. Voth @ U. Utah: Path-Integral Monte Carlo Simulations
- G. Scoles & K. Lehmann @ Princeton U.: Helium Clusters

* External Collaborators:

- T. Momose @ Kyoto U.: High Resolution IR Spectroscopy

* Summer Visiting Professors:

- R.J. Hinde @ U. Tennessee: Dopant-Induced IR Activity
- D. Anderson @ U. Wyoming: Dopant IR Absorptions

Propellant Performance Figures of Merit

Specific Impulse, I_{sp} :

$I_{sp} \equiv$ (total impulse / propellant weight)

$= g_0 \langle v_{exh} \rangle$ "seconds"

$$\propto \sqrt{\frac{\langle K.E. \rangle}{m}} \propto \sqrt{\frac{\Delta H}{m}} = \sqrt{\Delta H_{sp}}$$

Density, ρ :

higher density \Rightarrow smaller & lighter tanks

\Rightarrow less aerodynamic drag

\Rightarrow condensed phase propellants

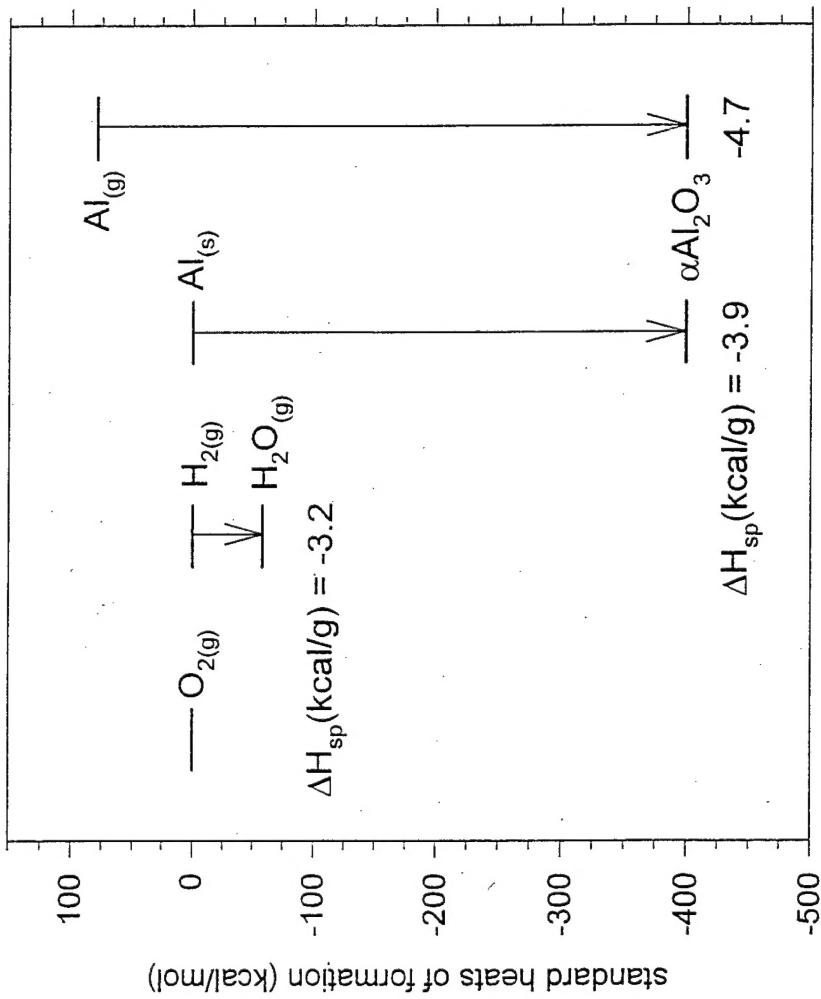
[G.P. Sutton, "Rocket Propulsion Elements" (Wiley, New York, 1992).]

“Revolutionary” vs. “Evolutionary” HEDM Concepts

- * “Revolutionary” means better than LOX/LH₂:
LOX/LH₂ $\Delta H_{sp} = 12.6 \text{ MJ/kg}$ (3.0 kcal/g)
HEDM Target: $\Delta H_{sp} > 15.0 \text{ MJ/kg}$ (3.6 kcal/g)
- * Early (c1990) Revolutionary HEDM Concepts:
 - tetrahydrogen (H₄)
 - metastable triplet helium (He* and He₂*)
 - spin-polarized atomic hydrogen (H↑)
 - high-spin species (⁵CO)
 - dications (AB⁺⁺, ABC⁺⁺)
- ▼ “non-metallics” (e.g. O₄/H₂, N₄, N₈, N₂₀) N₅⁺!
 - metallic hydrogen
 - metal atoms and clusters in solid H₂

Cryosolid Propellants Concept

Use cryogenic solid hydrogen as a “packaging material” to store energetic species such as metal atoms and clusters.



Atom Additive Payoffs (5 % molar)

Sea level specific impulse, I_{sp} , in seconds (% change)

$P_{\text{chamber}} = 1000 \text{ PSIA}$, $P_{\text{exhaust}} = 14.7 \text{ PSIA}$

Additive	in standard state <u>M(5%)/LOX/H₂</u>	as atoms <u>M(5%)/LOX/H₂</u>	monoprop. <u>M(5%)/H₂</u>
none	403		
C	381 (-5%)	515 (+28%)	515 (+28%)
B	407 (+1%)	508 (+26%)	465 (+15%)
Be	427 (+6%)	493 (+22%)	
Si	400 (-1%)	460 (+14%)	
Al	407 (+1%)	454 (+13%)	
H	403	430 (+7%)	380 (-6%)
Li	404	428 (+6%)	
Mg	400 (-1%)	416 (+3%)	

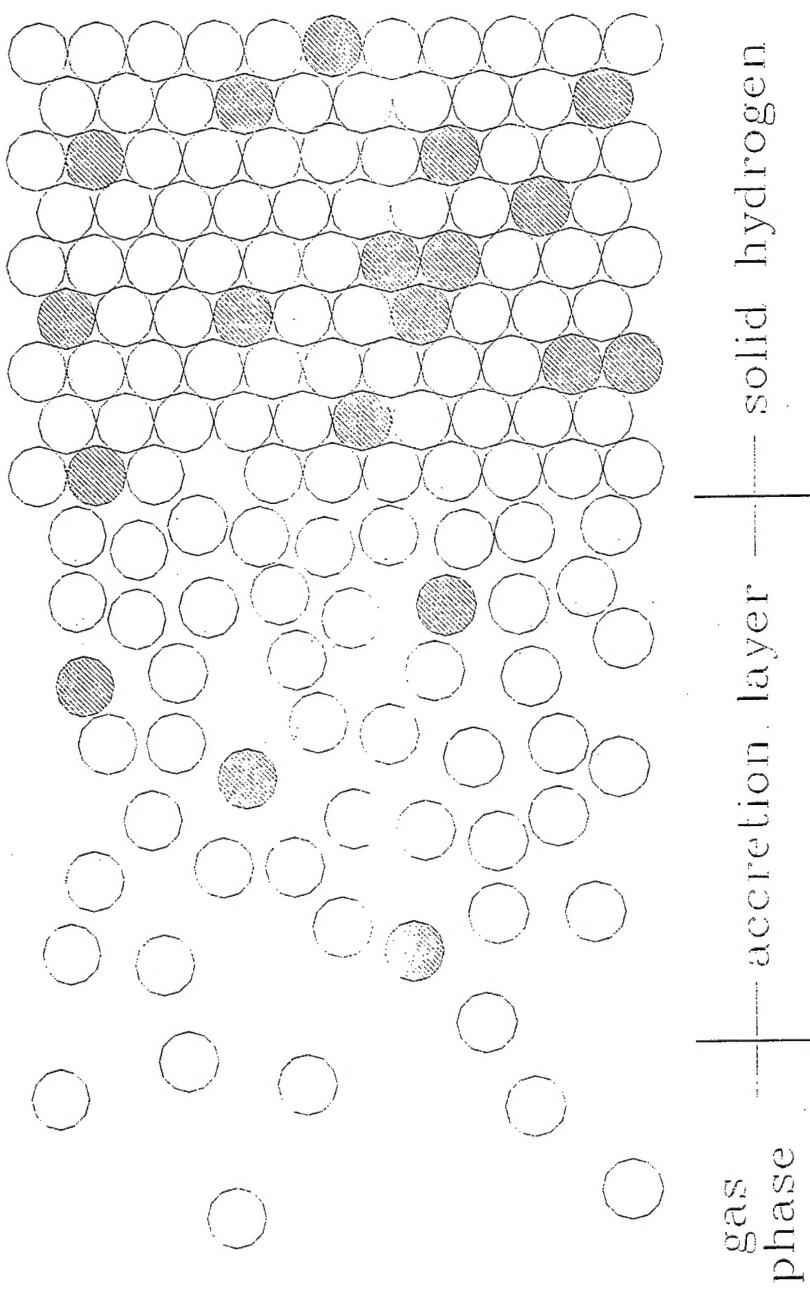
Cryosolid Propellants Objectives

- * Make solid hydrogen samples (any size) containing 5% molar concentration of trapped energetic additives.
- * Measure absolute concentrations of energetic species.
- * Scale-up samples; produce $\sim 1 \text{ cm}^3$ samples in our lab.

Example: 5% Al/pH₂, V = 1 cm³
assume each Al atom replaces one H₂ molecule
⇒ 58 mg Al / 83 mg H₂ (*see display item*)
∴ ρ = 0.142 g/cm³ (+100%)

Cryosolid Propellants Approach

- * Rapid vapor deposition of metal atom vapor and pre-cooled parahydrogen gas onto a liquid helium cooled substrate in vacuum.



Dopant Reactions within solid pH₂

- * ideally:
$$X + pH_2 \xrightarrow{T=2K} X/pH_2$$
 isolated atoms
- * in practice:
$$X + X + M \rightarrow X_2 + M$$

$$\rightarrow X_n$$
 recombination
- $$X + H_2 + M \rightarrow HX + H + M$$

$$\rightarrow H_nX + M$$
 reaction
- $$X_n + H_2 + M \rightarrow HX_n + H + M$$

$$\rightarrow H_mX_n + M$$
 both

The Perils of Calorimetry

TABLE IX
CONCENTRATIONS OF FREE RADICALS REPORTED

Radical	Matrix	Mole per cent radicals	Method of production and estimate ^a	Reference
O	O ₂	4-20	Gas, cal	Minkoff <i>et al.</i> (1959)
		<3	Gas, IR	Harvey and Bass (1958)
		~1	Gas, cal	Broida and Lutes (1956)
	Ca(OH) ₂	0.6	γ, ESR	R. Livingston ^b
	N ₂	4	Gas, cal	Minkoff <i>et al.</i> (1959)
		0.2	Gas, cal	Broida and Lutes (1956)
		0.03	γ, ESR	Wall <i>et al.</i> (1959b)
		>0.03	Gas, cal	Fontana (1958)
		0.01-0.04	Gas, MS	Fontana ^c
		0.2	γ, ESR	Matheson and Smaller (1955)
OH(?)	HCOOH	0.14	γ, ESR	Wall <i>et al.</i> (1959a)
	CH ₄	0.1	γ, ESR	Wall <i>et al.</i> (1959a)
	NH ₃	0.1	Gas, ESR	Cole and Harding (1958)
	HClO ₄ -H ₂ O	0.1	γ, ESR	Livingston <i>et al.</i> (1955)
	H ₂ O	0.01	γ, ESR	Matheson and Smaller (1955)
	H, NH ₂ (?)	0.01	UV, ESR	D. Ingram ^b
	ROH	~0.01	UV, ESR	Wall <i>et al.</i> (1959a)
	H ₂	0.0006	γ, ESR	
	H			

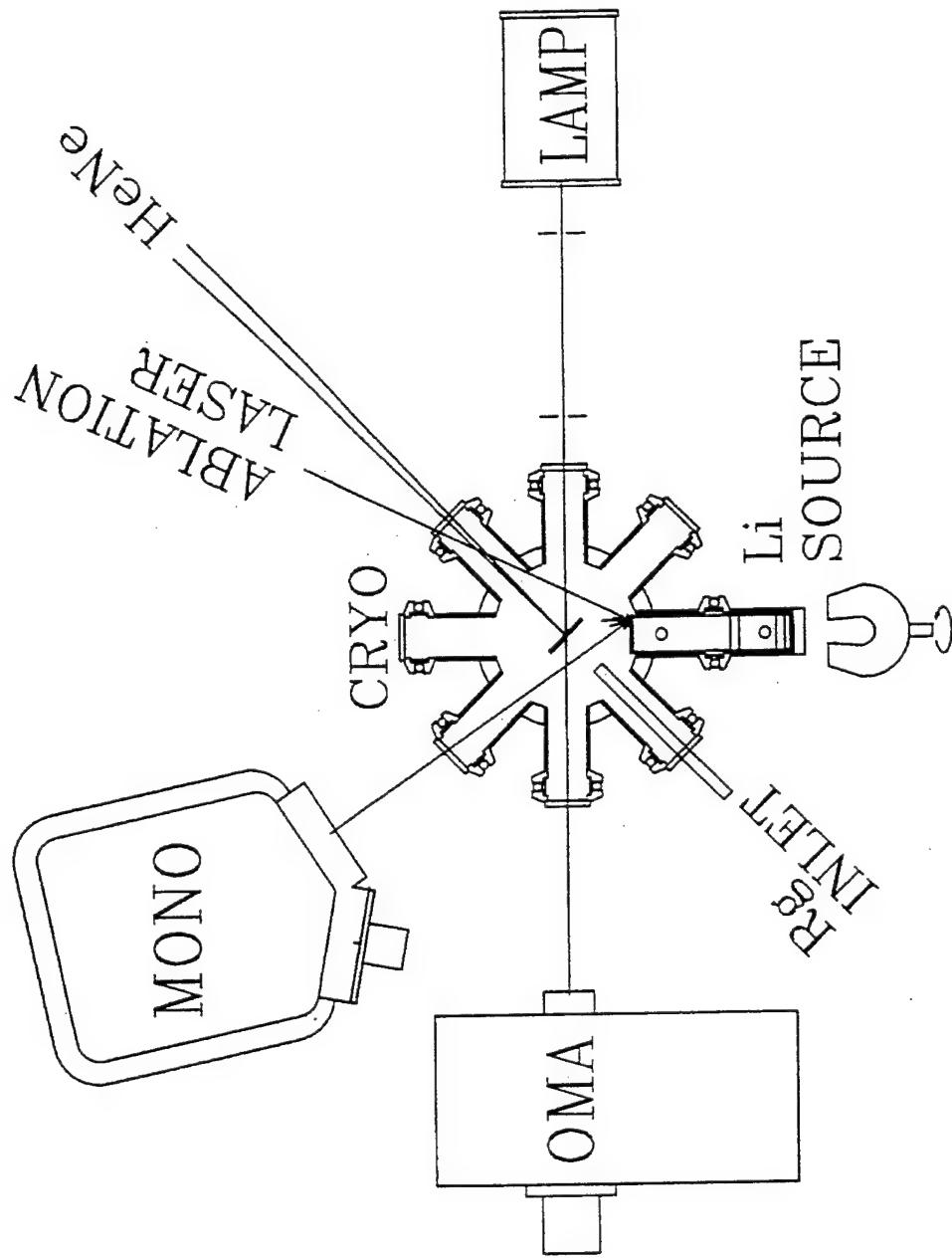
^a Abbreviations: gas = rapid condensation of gaseous radicals; γ = gamma ray; *in situ* production; UV = photolytic *in situ* production; IR = infrared analysis; cal = calorimetry; MS = magnetic susceptibility.

^b Private communication.

^c Fontana, B. J. (1959). *J. Chem. Phys.* **31**, 148.

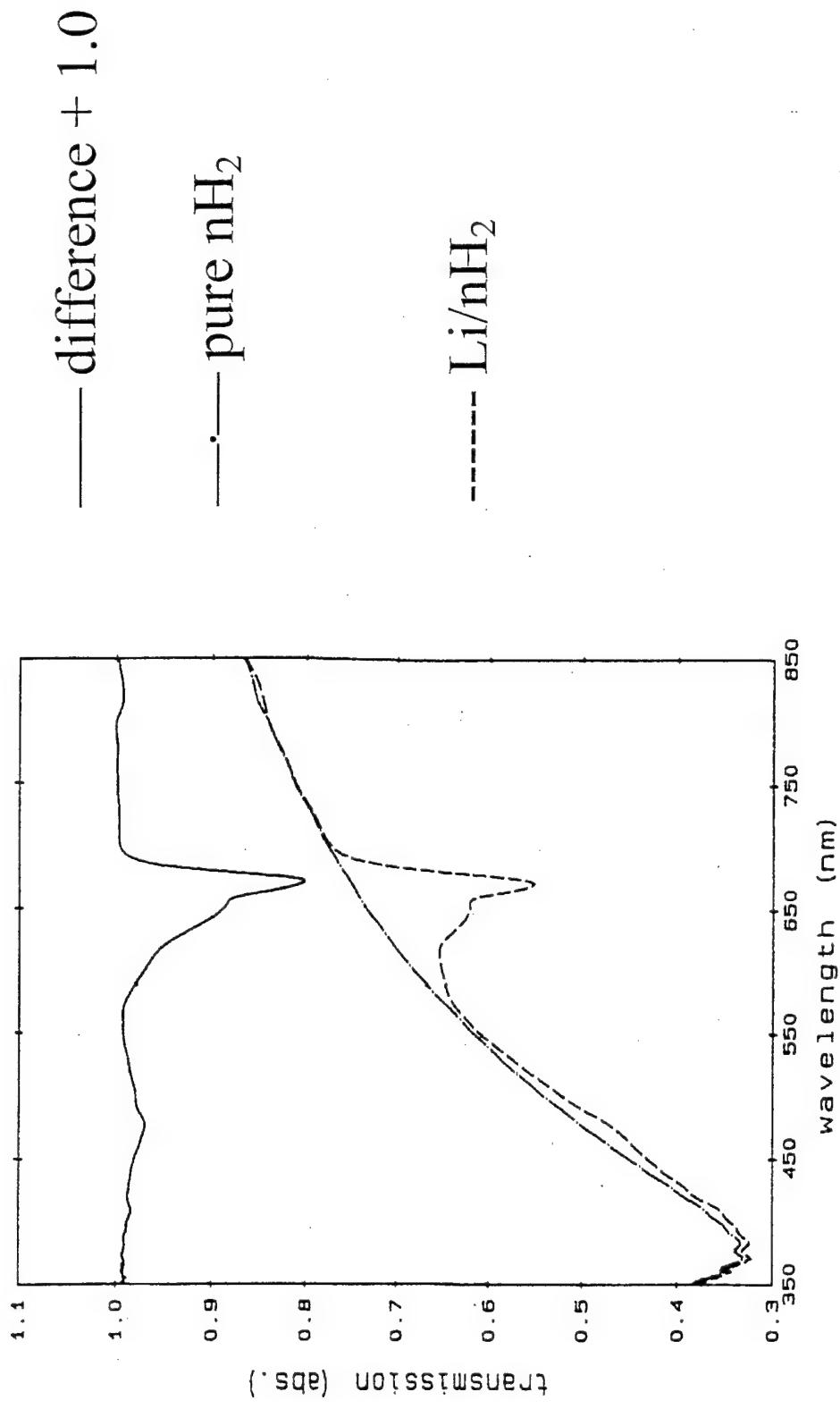
[A.M. Bass and H.P. Broida, "Formation and Trapping of Free Radicals" (Academic, New York, 1960).]

Experimental Diagram (c1993)



M.E. Fajardo, J. Chem. Phys. **98**, 110 (1993).

Transmission Spectrum of Li/nH₂, d ≈ 10 μ



Optical Scattering in Solid Hydrogen

Crystal Growing and Quality (p. 81)

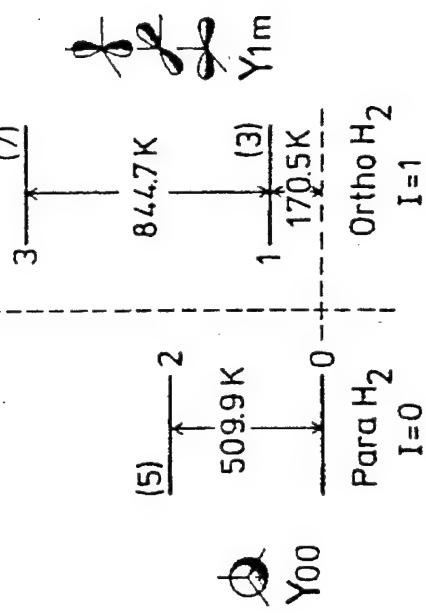
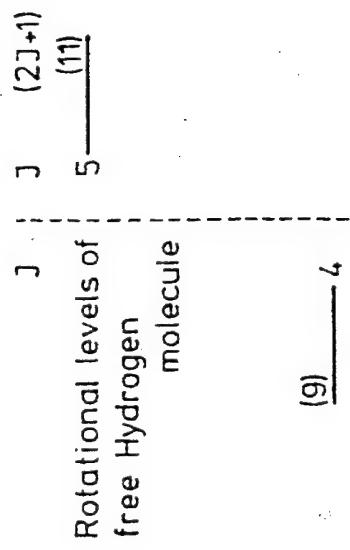
“There is a considerable art to growing hydrogen crystals of high quality. Good crystals are always grown slowly from the melt; a rapid freeze from the gas produces snow.”

Crystallite Light Scattering (p. 83)

“The reason that a good hydrogen crystal is so hard to see is its low refractive index...an estimated 1.16!

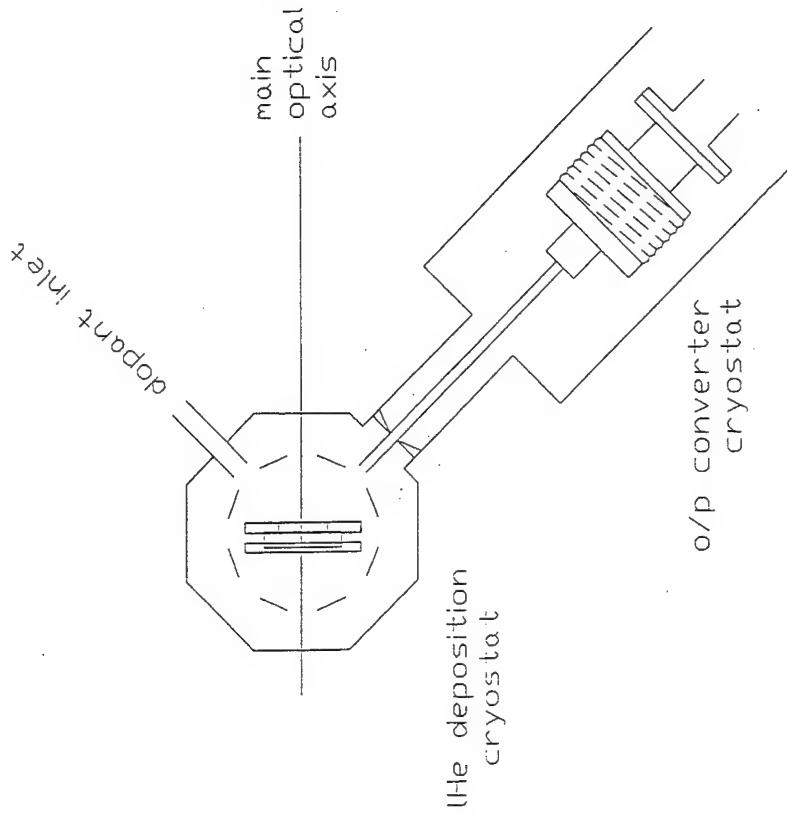
Yet a 1 mm-thick layer of hydrogen crystallites can be a completely opaque brown-black.”

ortho- and para-hydrogen



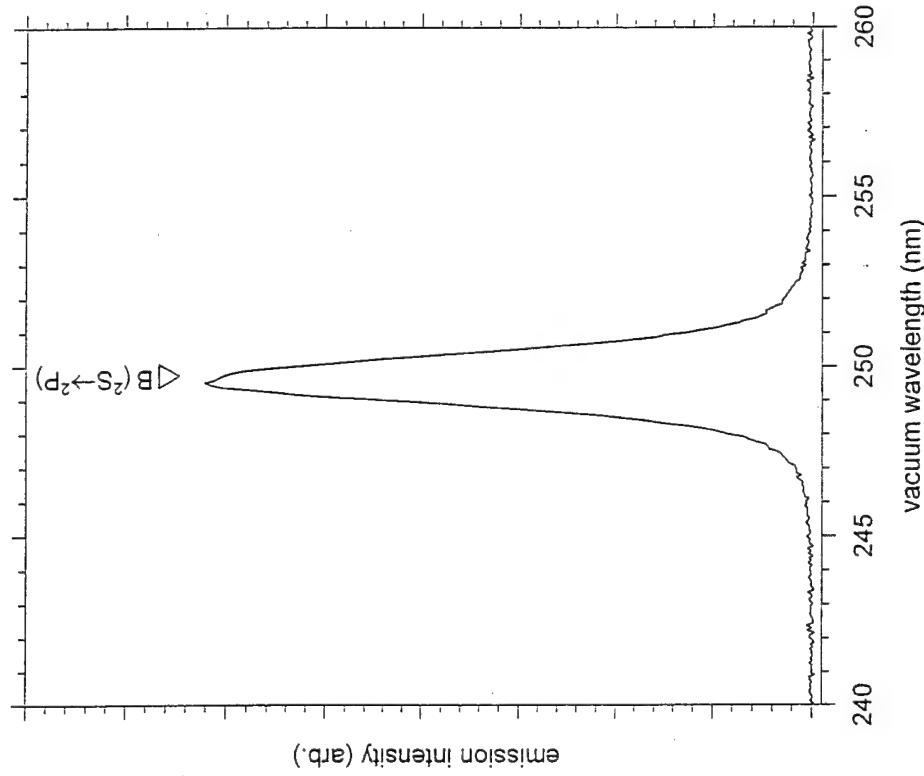
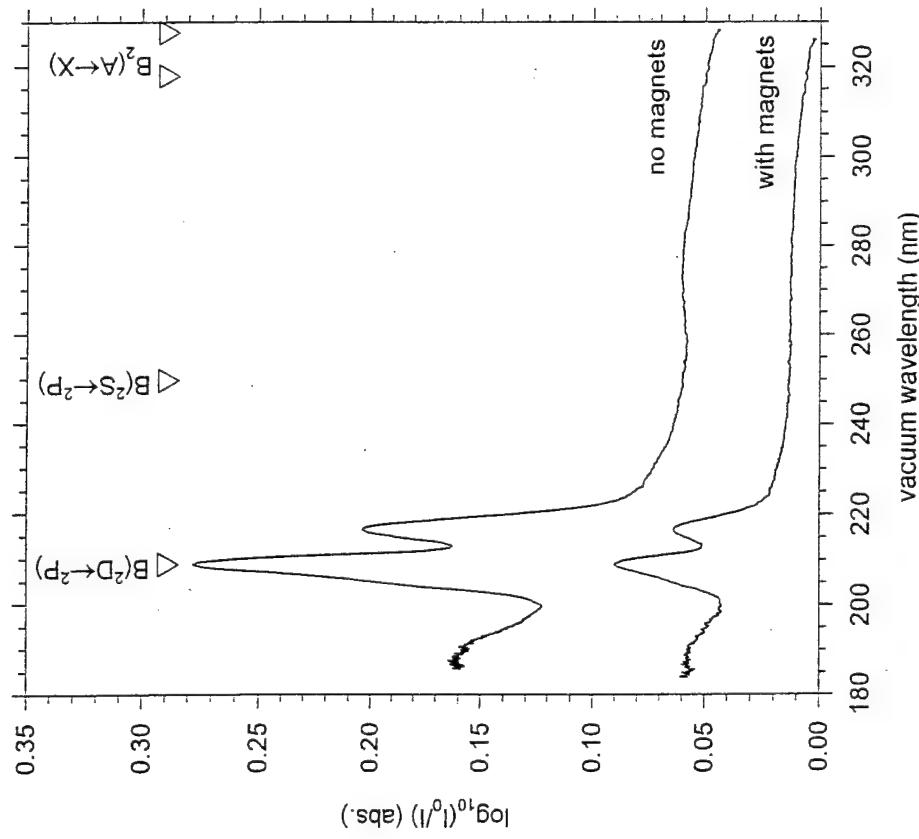
[I.F. Silvera, Rev. Mod. Phys. 52, 393 (1980)]

Rapid Vapor Deposition of Gram-Scale Optically Transparent pH₂ Solids (c1997)



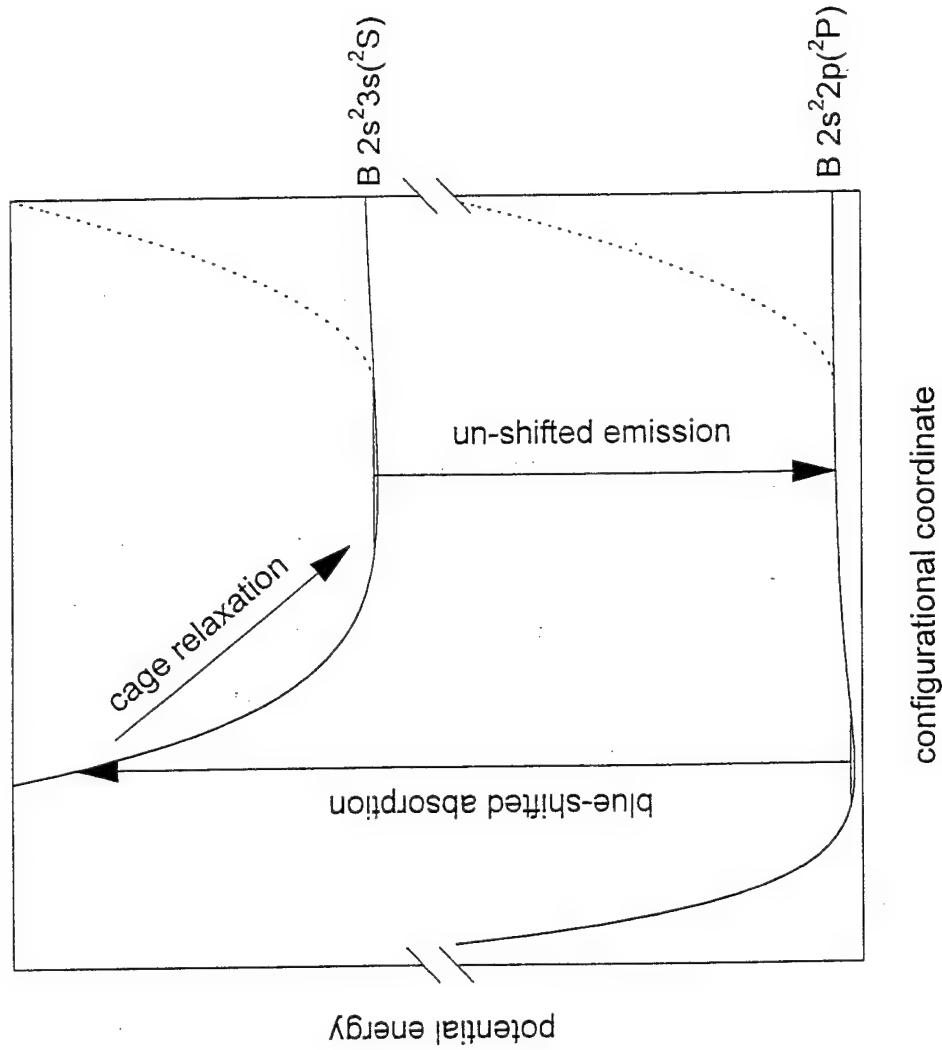
M.E. Fajardo and S. Tam, J. Chem. Phys. **108**, 4237 (1998).
S. Tam and M.E. Fajardo, Rev. Sci. Instrum. **70**, 1926 (1999).

Electronic Spectroscopy of B/pH₂ (d≈2 mm)



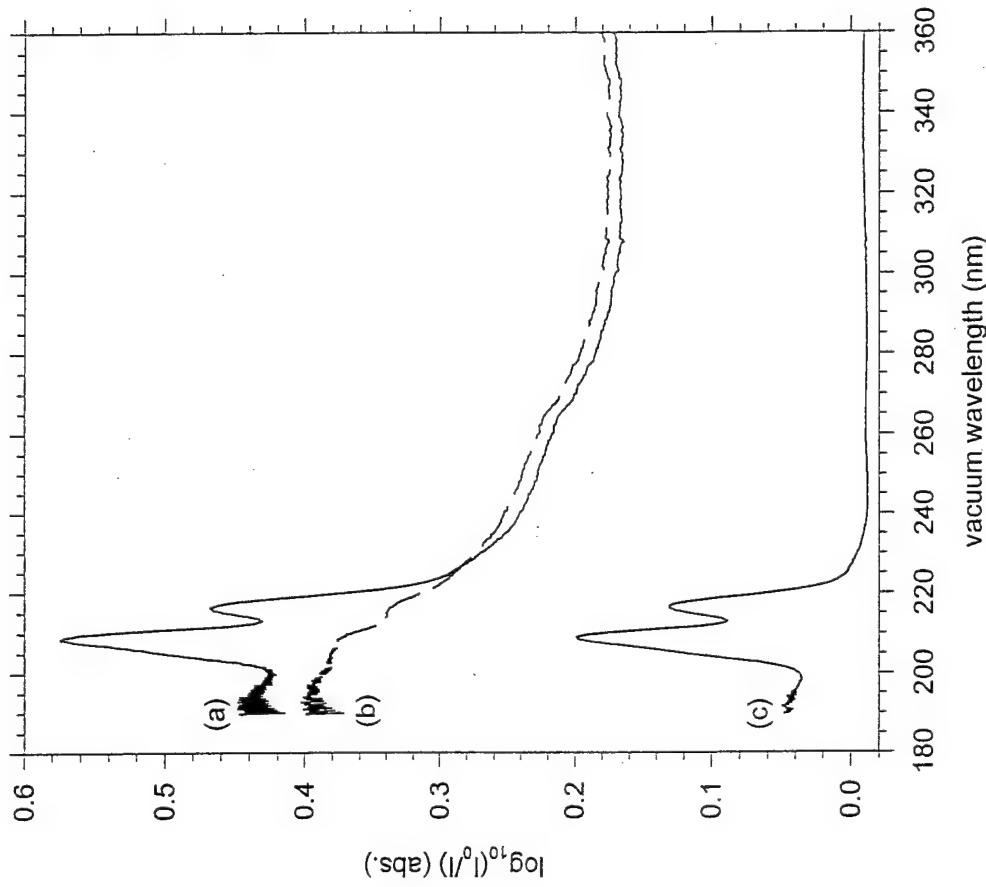
S. Tam, M. Macler, M.E. DeRose, and M.E. Fajardo, J. Chem. Phys. **113**, 9067 (2000).
[J.R. Krumrine, S. Jang, G.A. Voth, and M.H. Alexander, J. Chem. Phys. **113**, 9079 (2000)]

Photodynamics Cartoon for B/pH₂



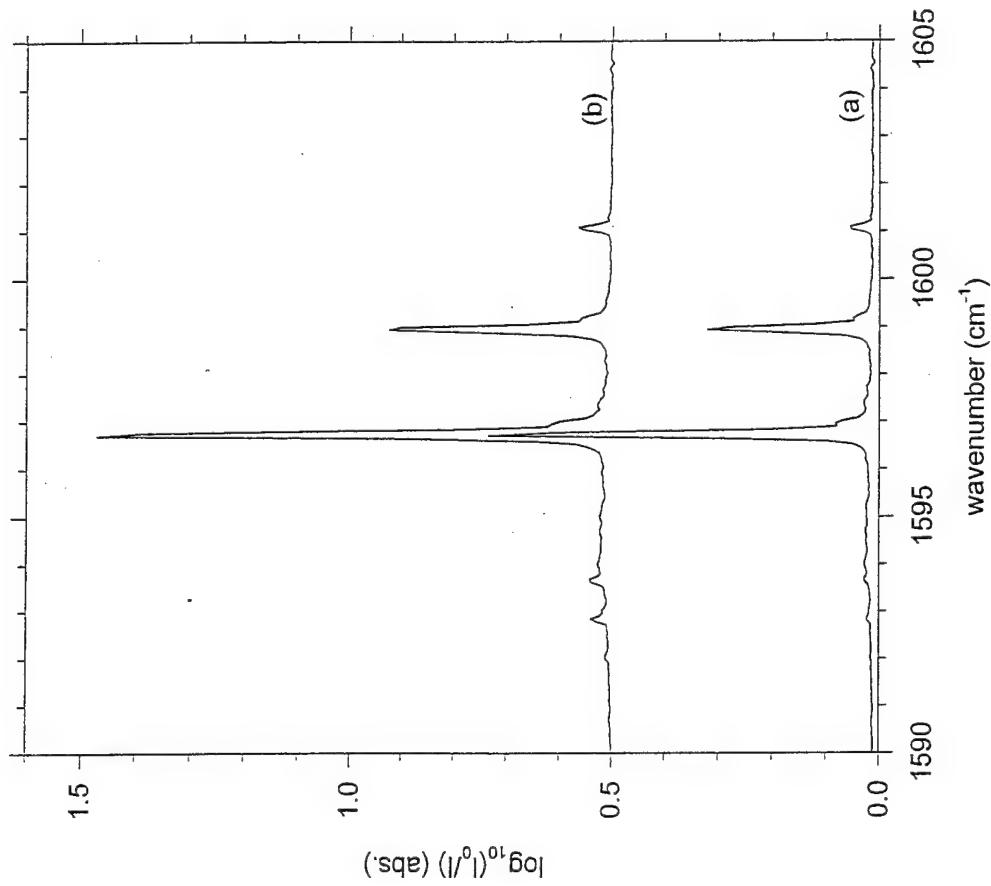
S. Tam, M. Macler, M.E. DeRose, and M.E. Fajardo, J. Chem. Phys. **113**, 9067 (2000).

Photobleaching of B/pH₂ Absorptions



S. Tam, M. Macler, M.E. DeRose, and M.E. Fajardo, J. Chem. Phys. **113**, 9067 (2000).

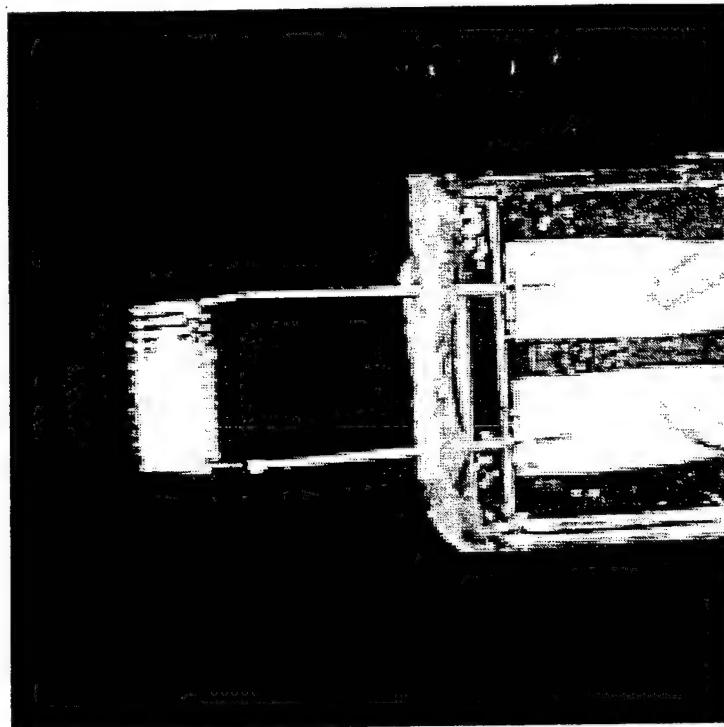
IR Absorption Spectra of B₂H₆/pH₂



S. Tam, M. Macler, M.E. DeRose, and M.E. Fajardo, J. Chem. Phys. **113**, 9067 (2000).

Requirement for High Flux HEDM Sources

- * A 5 % doping level, and a sample growth rate of 1 mm/hour, require a flux of $\Phi_{\text{HEDM}} \approx 3 \times 10^{16} \text{ #}/\text{cm}^2\text{-s}$ at the deposition substrate. For Al atoms, this translates to a mass flux of $5.8 \text{ mg}/\text{cm}^2\text{-hour}$, delivered to the deposition substrate.
- * Began FY00 using miniature tungsten filament evaporation sources based on our FY99 effort to produce thermal B atoms.
- * Total mass loadings of Al metal were $\sim 10 \text{ mg}$, just enough to detect trapped Al atoms in Ar; $\Phi_{\text{Al}} \sim 10^{11} \text{ #}/\text{cm}^2\text{-s} @ R = 5 \text{ cm}$.



High Flux HEDM Sources

- * Purchased commercial Al evaporator; PBN crucible holds ≈ 10 g Al in horizontal orientation.
- * $T_{\max} = 1200^{\circ}\text{C} \Rightarrow P_{\text{vap}}(\text{Al}) \approx 8 \times 10^{-3}$ torr $\Rightarrow \Phi_{\text{Al}} \approx 10^{18} \text{ #}/\text{cm}^2\text{-s}$

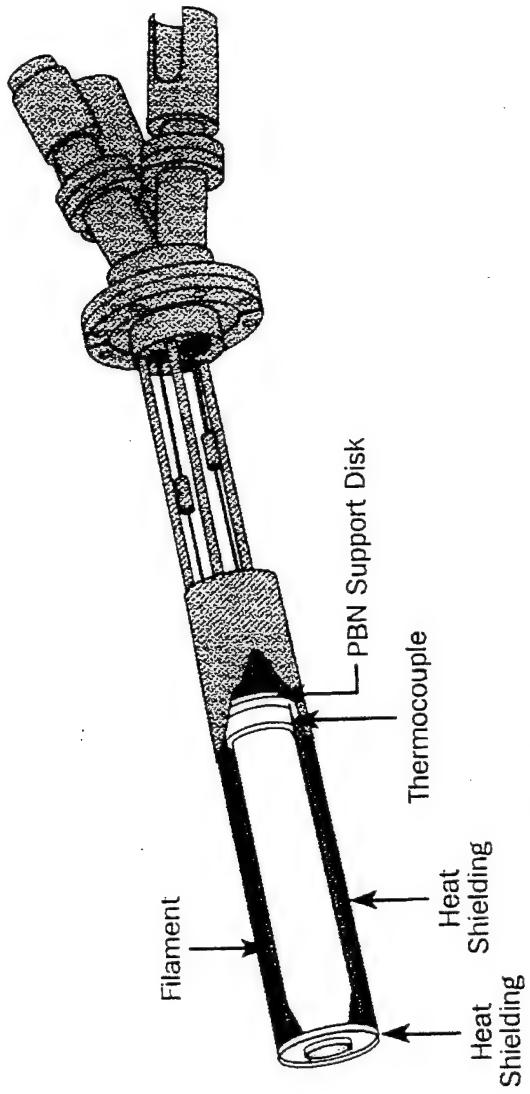
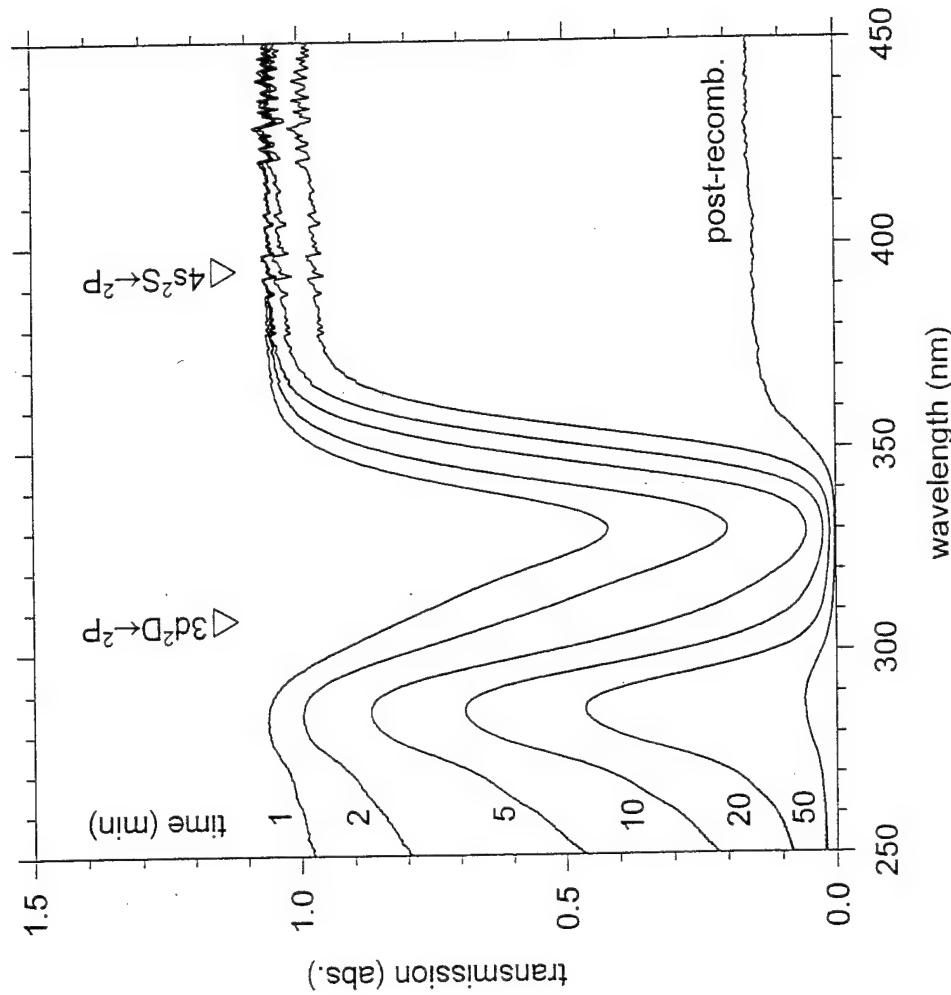


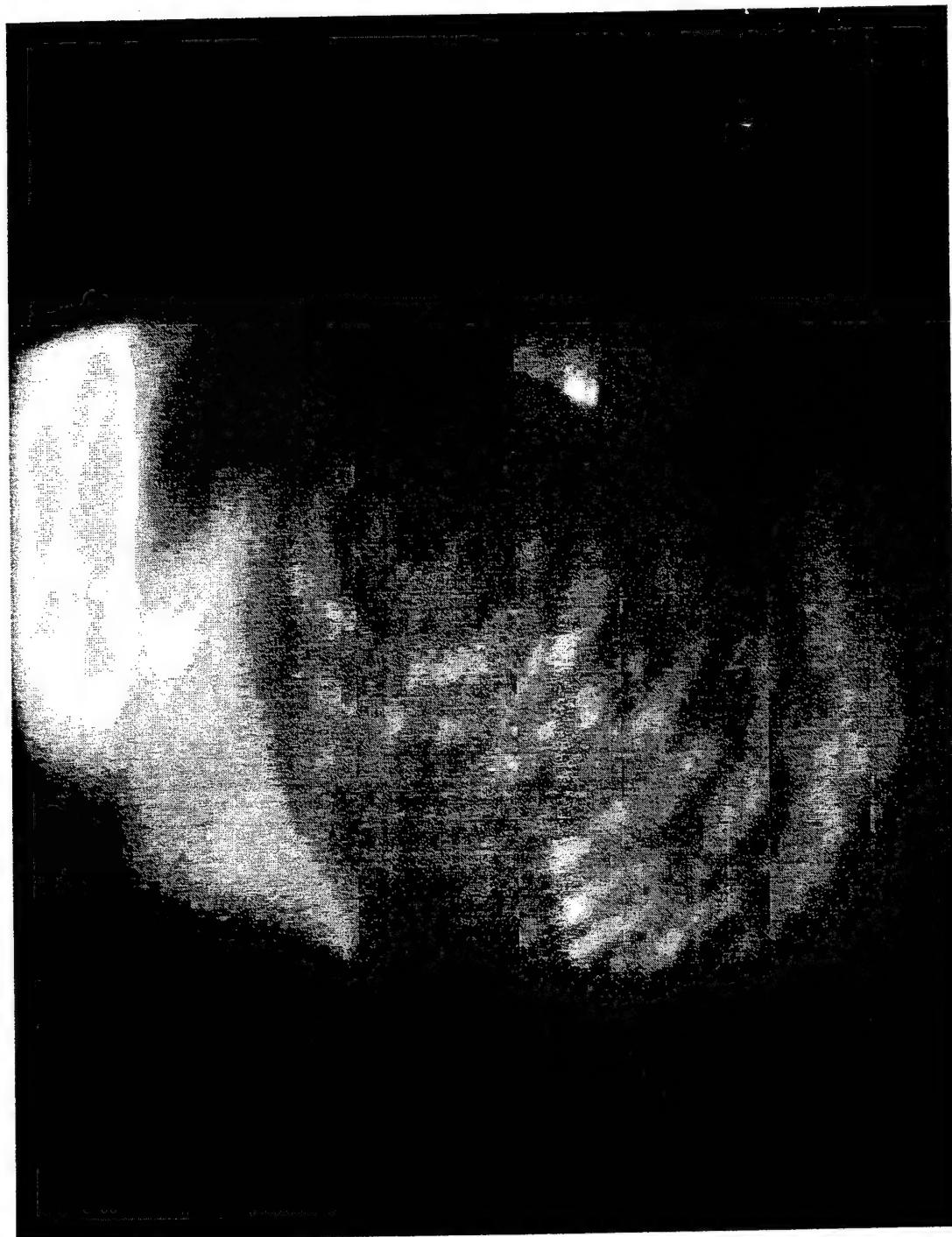
Figure I-3: Schematic of the EPI SUMO™ Effusion Cell.

UV Spectroscopy Al/pH₂

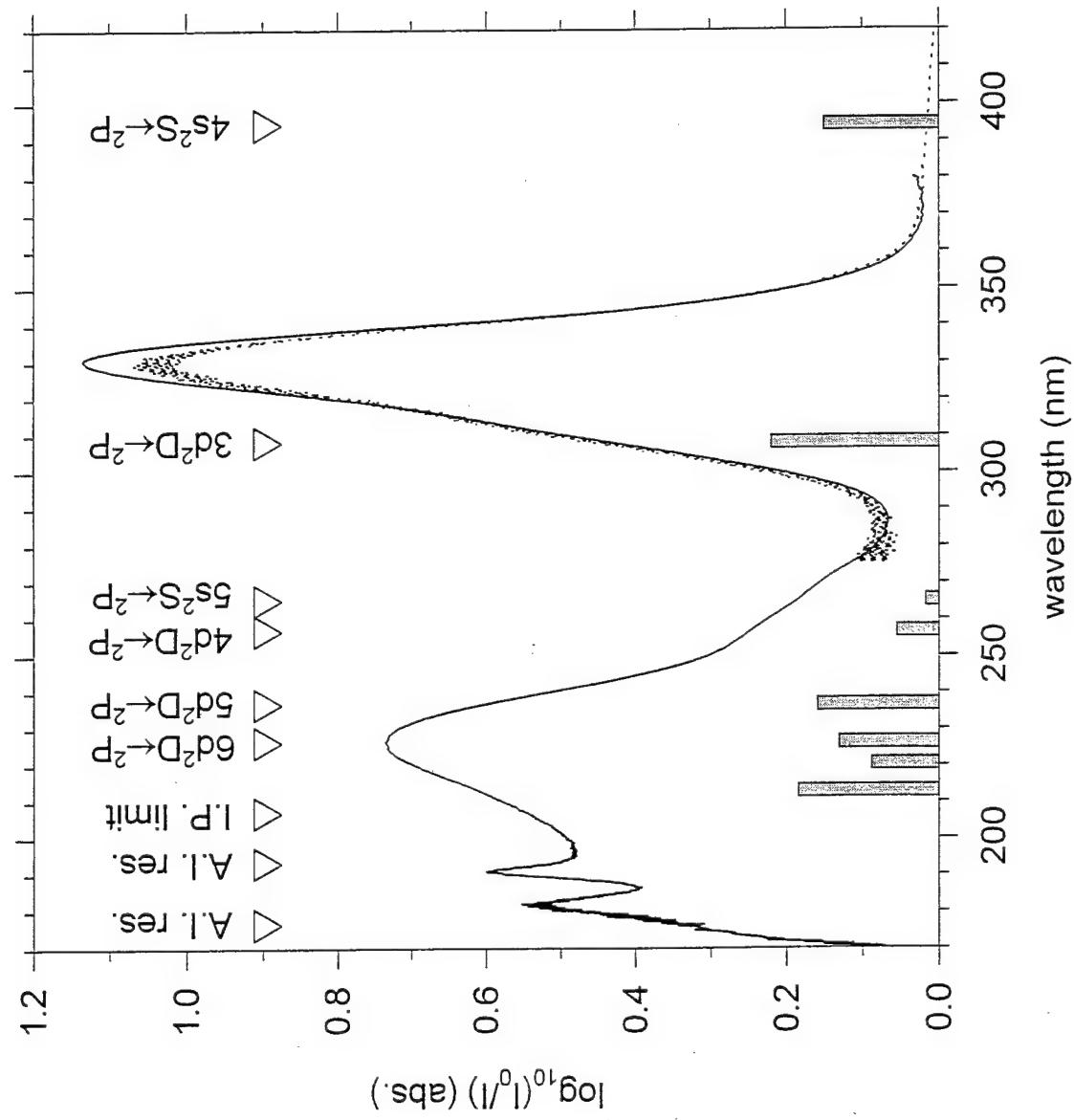
- * Al atom UV absorption saturates at high column densities.



Recombination/reaction in Al/pH₂



Assignment of Al/pH₂ UV Absorptions



High Res. IIR Spectroscopy in Solid pH₂

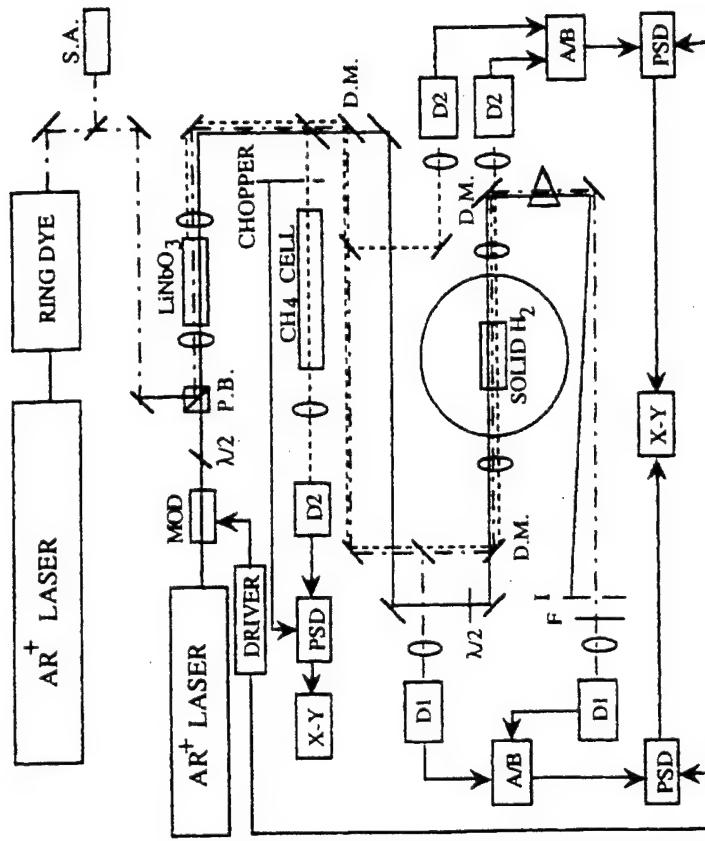
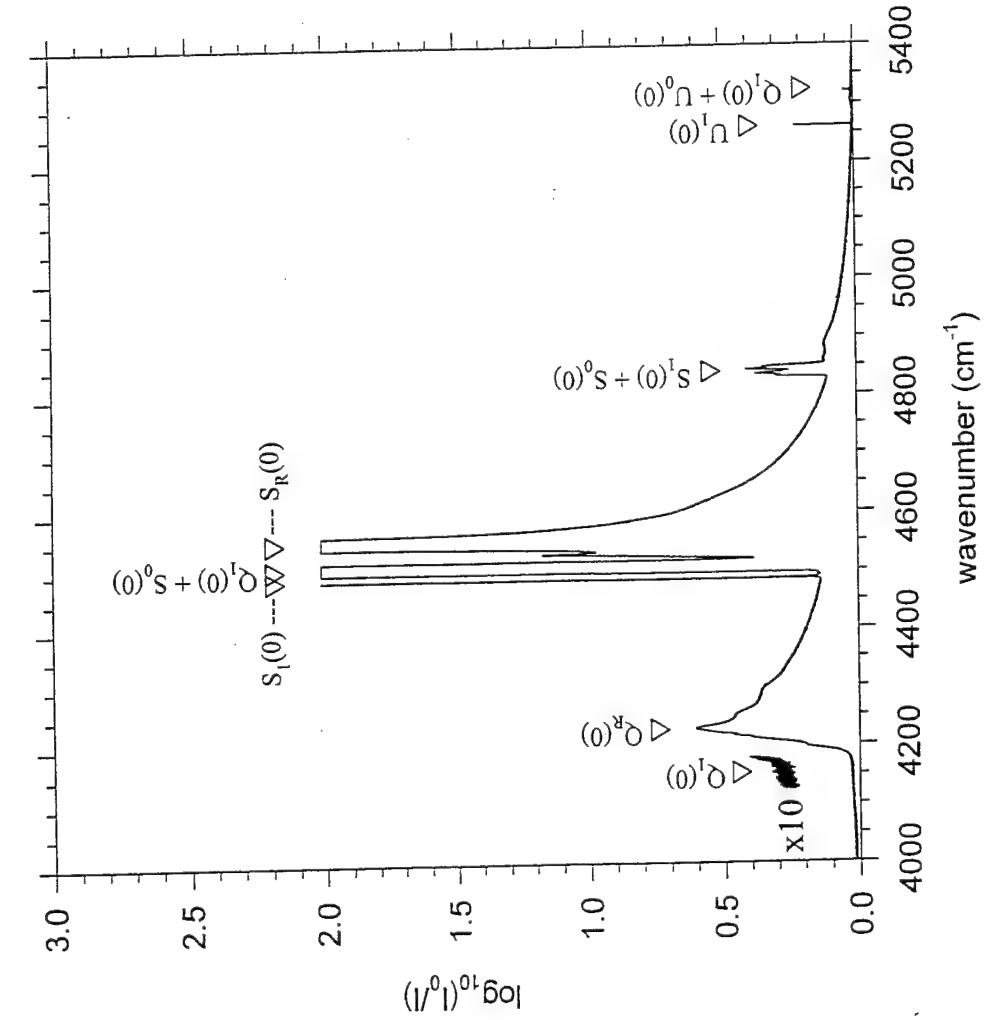


FIG. 1. Apparatus for the simultaneous spectroscopy of the infrared and Raman transitions. The nonlinearity of LiNbO_3 is used for the former and that of solid H_2 is used for the latter. D.M., dichroic mirror; S. A., spectrum analyzer; P. B., polarizer beamsplitter.

IR Absorption of 6 mm Thick pH₂ Solid



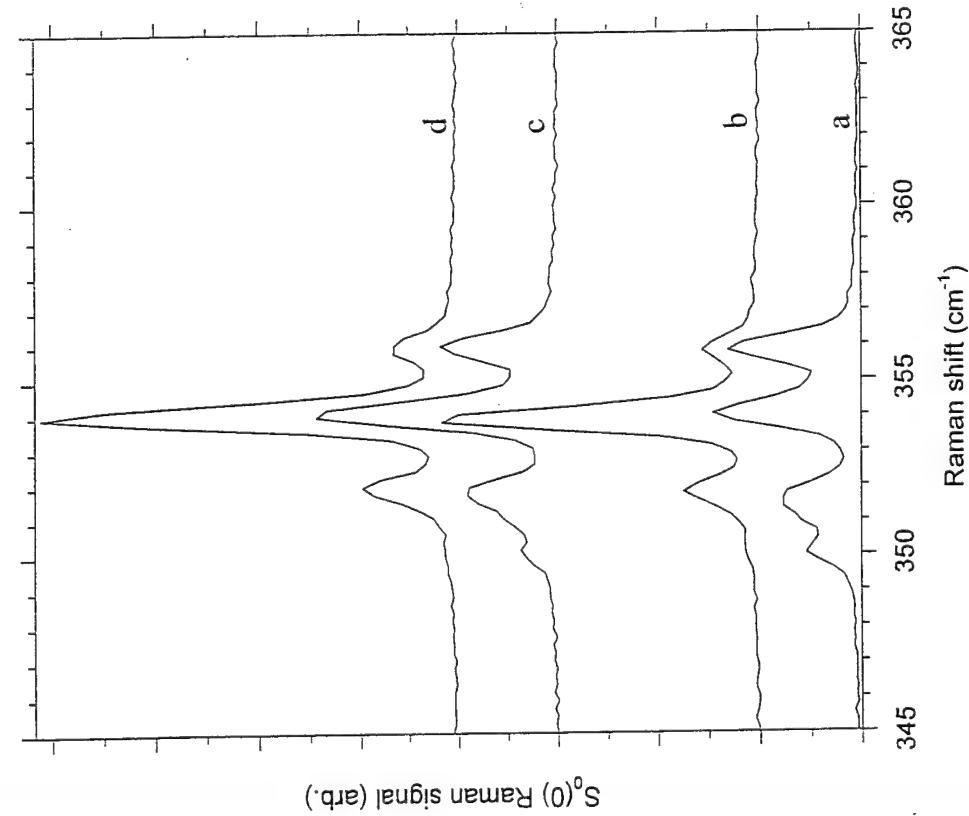
Non-observation of the $\text{Q}_1(0)$ transition demonstrates the absence of oH_2 impurities, and that the microscopic structure is not amorphous or porous.

Observation of $\text{S}_1(0)$ transition demonstrates the absence of inversion symmetry for some H_2 molecular environments.

[van Kranendonk and Gush, Phys. Lett. 1, 22 (1962)]

M.E. Fajardo and S. Tam, J. Chem. Phys. **108**, 4237 (1998).

Raman Spectra of pH₂ Solids



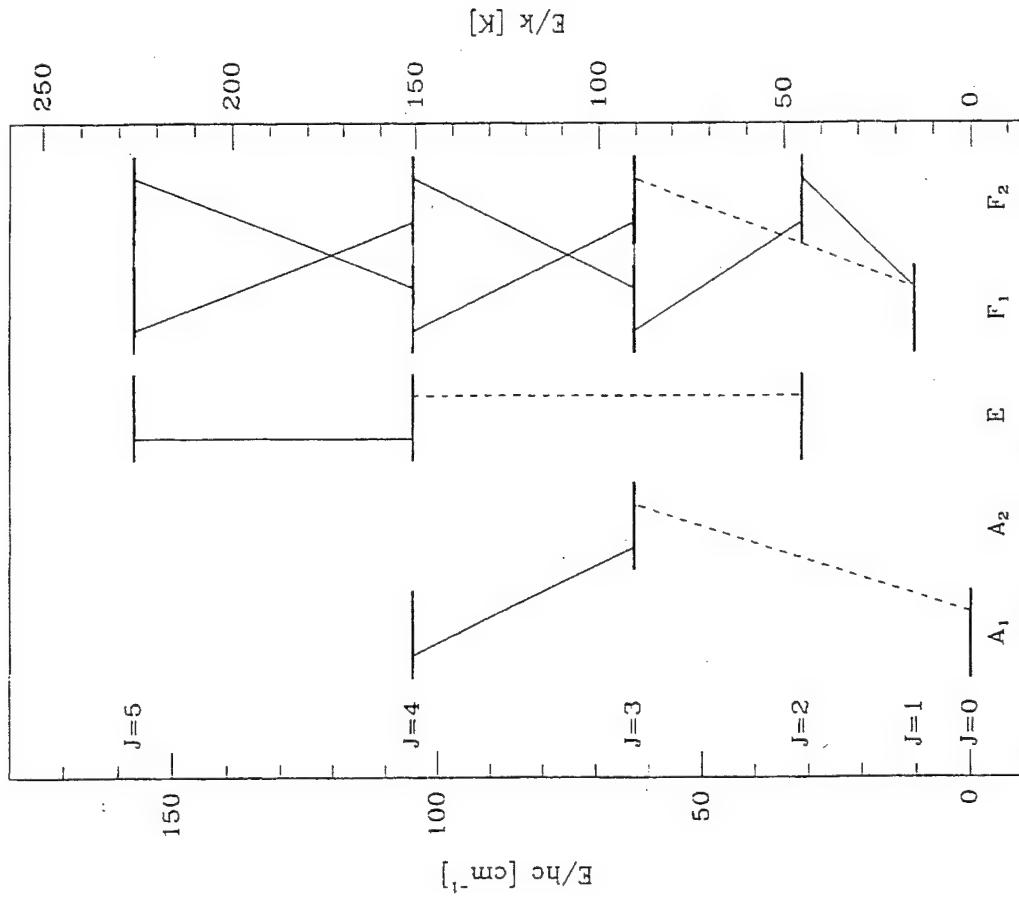
Mixed hcp/fcc as-deposited structure, anneals to hcp; compare with: [G.W. Collins, et al., Phys. Rev. B **53**, 102 (1996)].

(d) sample in (c) warmed to 4.5 K.
(c) 4.5 mm sample as deposited at 3.3 K ($\Phi = 290 \text{ mmol/hr}$).

(b) sample in (a) warmed to 4.5 K.
(a) 6 mm sample as deposited at 3.1 K ($\Phi = 200 \text{ mmol/hr}$).

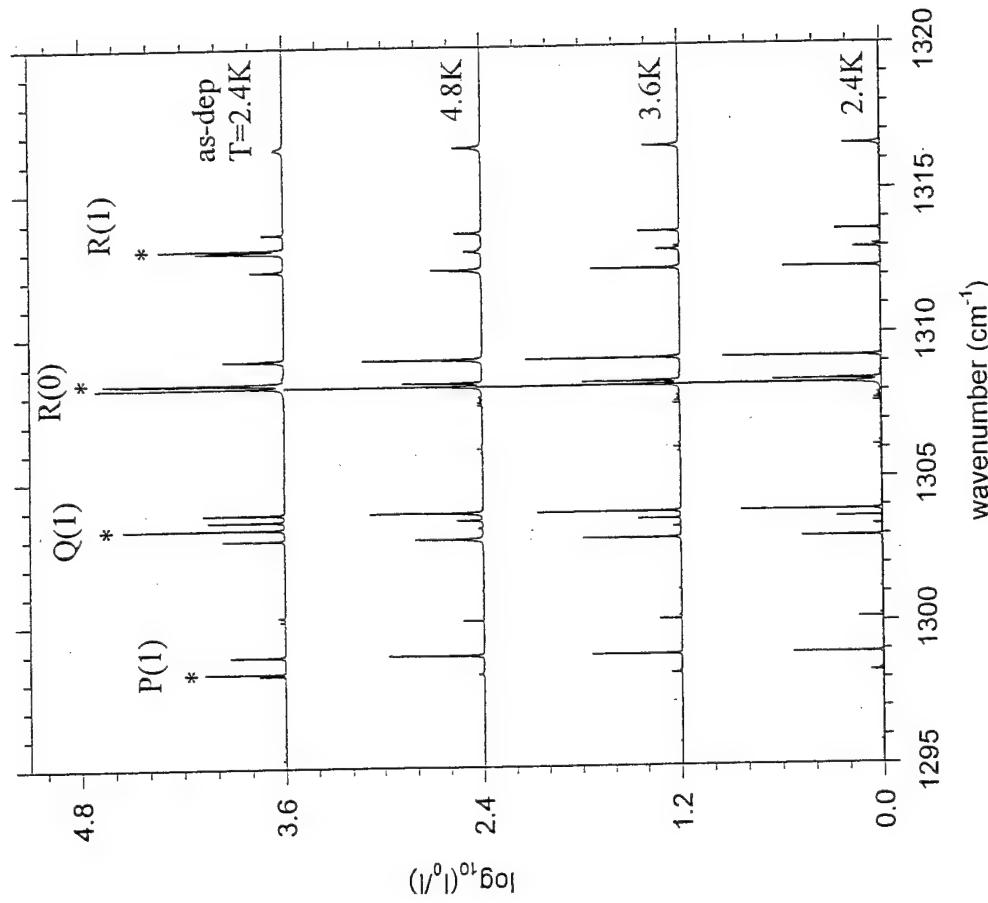
M.E. Fajardo and S. Tam,
J. Chem. Phys. **108**, 4237 (1998).

CH₄ Nuclear Spin Modifications



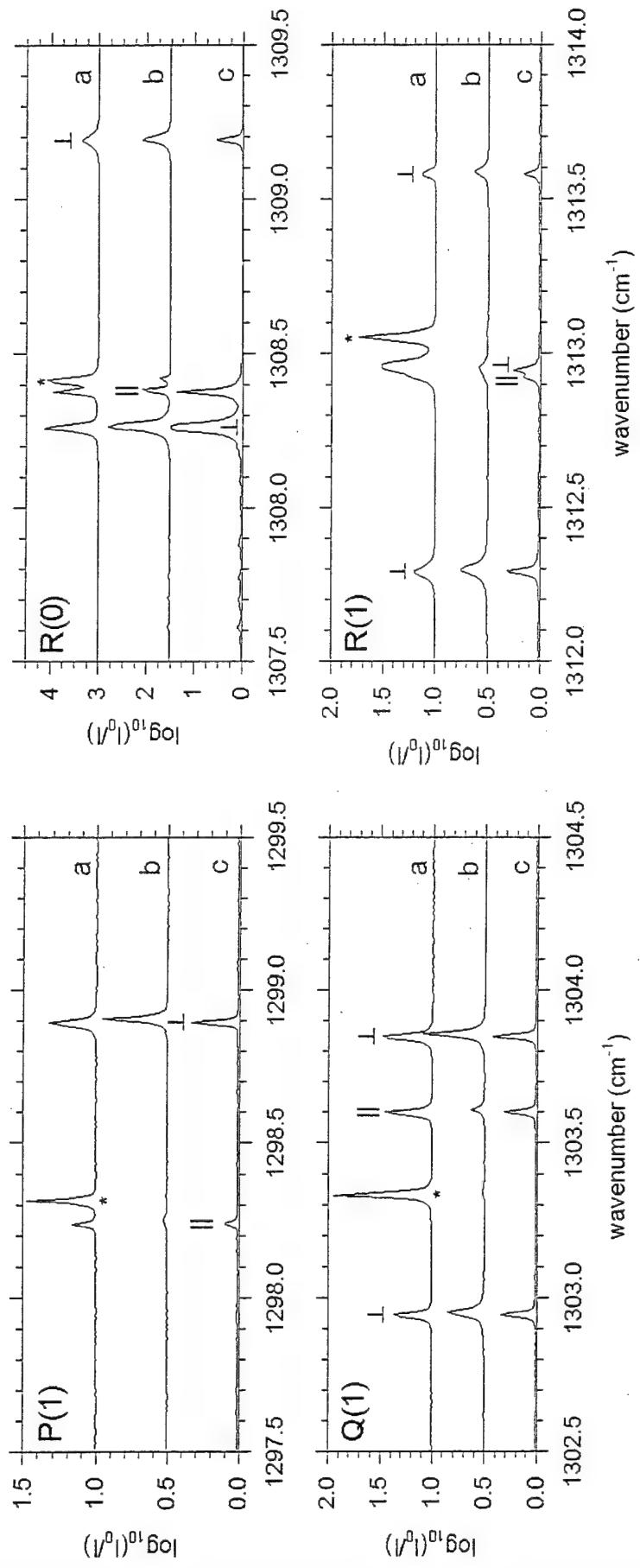
[M. Hepp, G. Winnewisser, and K.M.T. Yamada, J. Mol. Spectr. **164**, 311 (1994)]

ν_4 CH_4/pH_2 IR Absorptions (res = 0.01 cm^{-1})



S. Tam, M.E. Fajardo, H. Katsuki, H. Hoshina, T. Wakabayashi, and T. Momose, J. Chem. Phys. **111**, 4191 (1999).

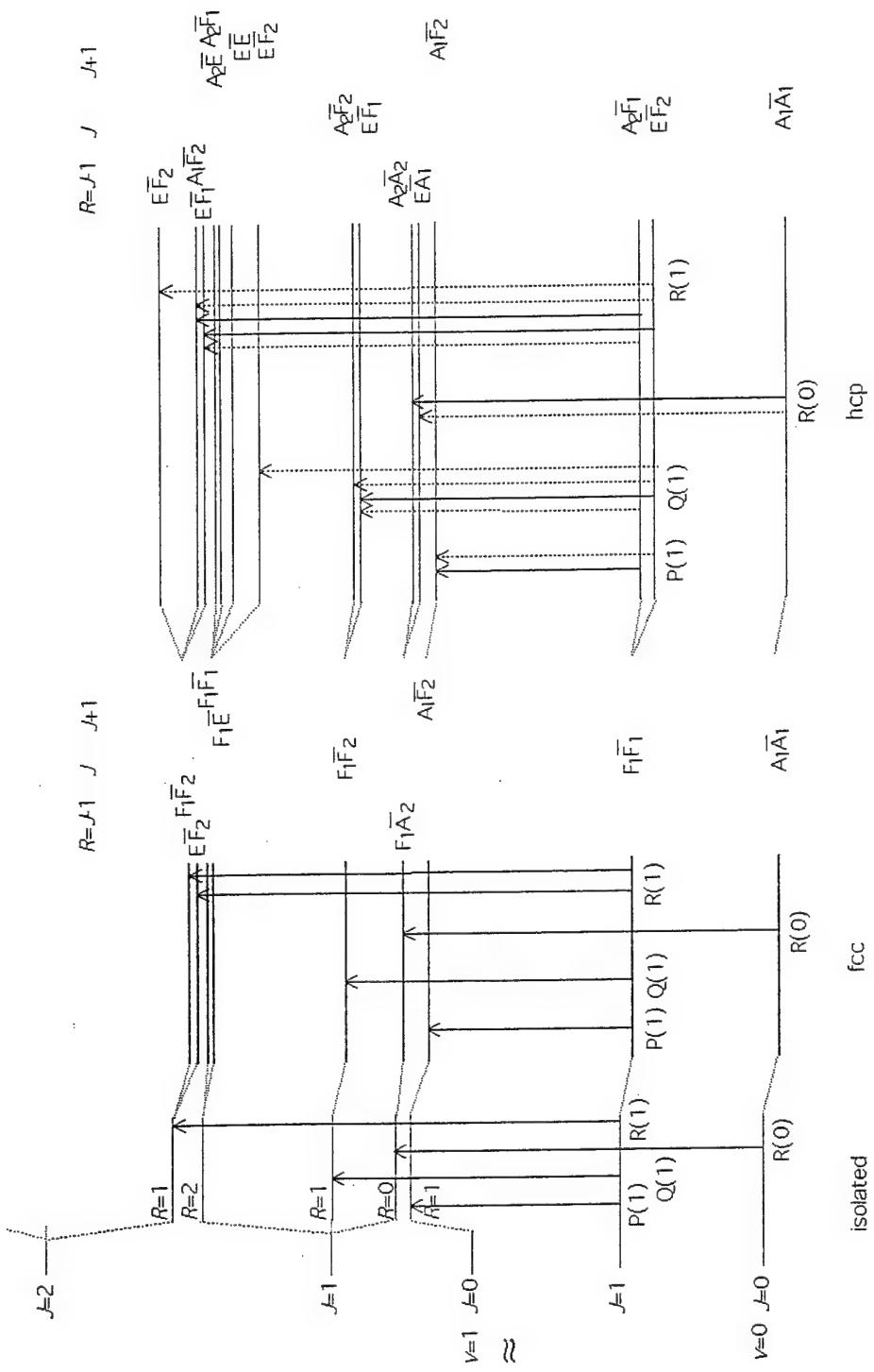
ν_4 CH_4/pH_2 IR Absorptions



- (a) Rapid Vapor Deposited sample: as-deposited at 2.4 K
- (b) Rapid Vapor Deposited sample: annealed to 4.8 K
- (c) Enclosed Cell Condensed sample: cooled to 4.8 K

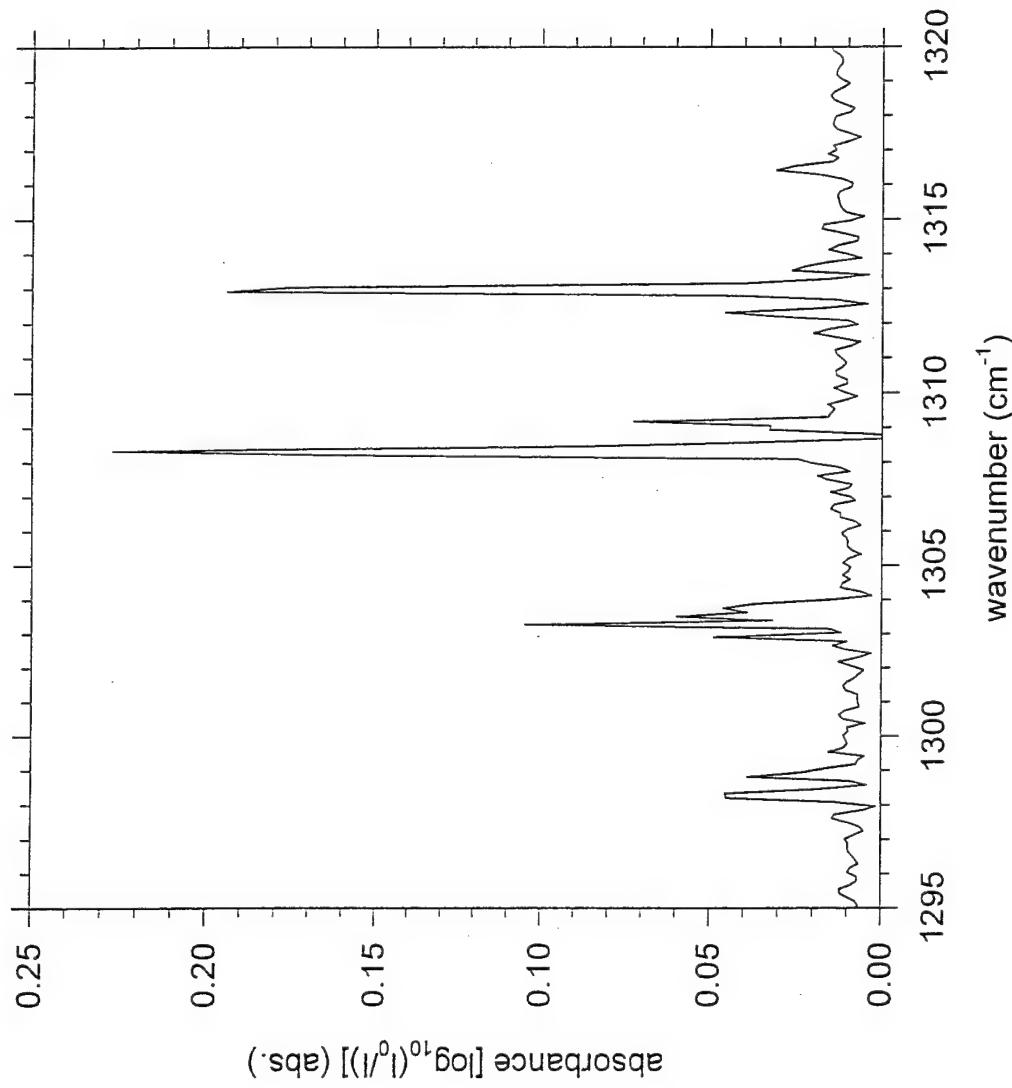
ν_4 CH_4/pH_2 Energy Levels

Department of Chemistry, University of Wyoming, Laramie, WY, 20 April 2001

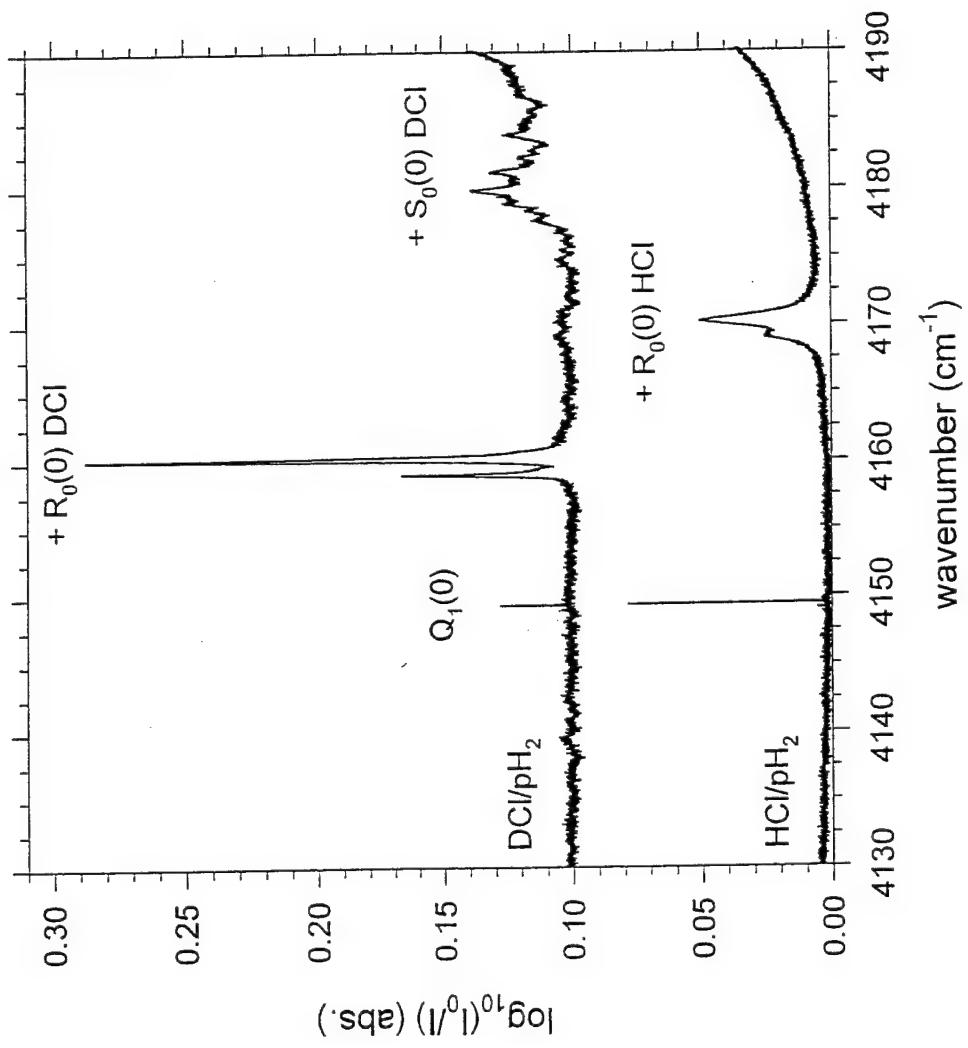


S. Tam, M.E. Fajardo, H. Katsuki, H. Hoshina, T. Wakabayashi, and T. Momose, *J. Chem. Phys.* **111**, 4191 (1999).

CH₄/pH₂ from Laser Ablation of Graphite

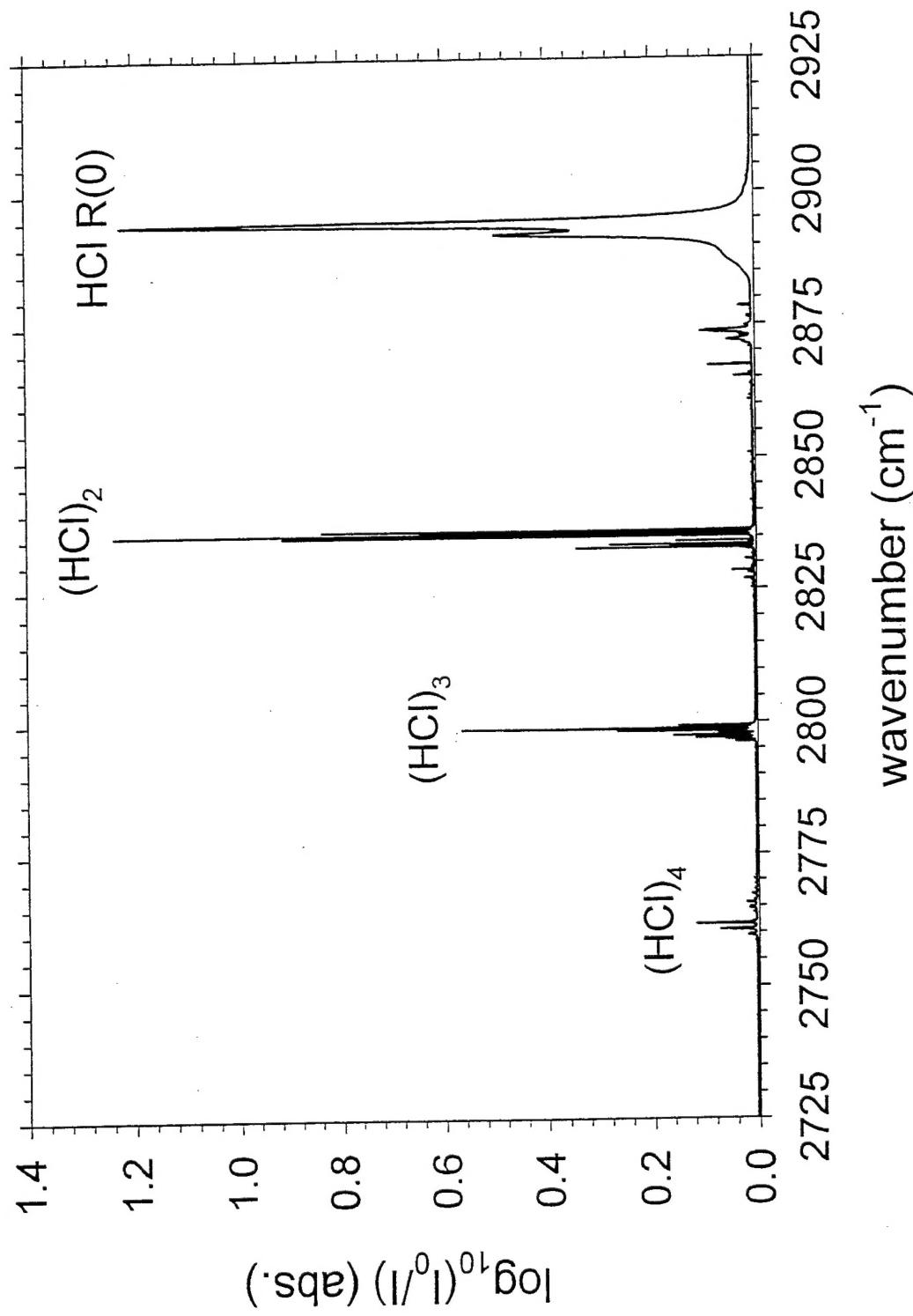


Co-operative IR absorptions



analysis in collaboration with D.T. Anderson, U. Wyoming and R.J. Hinde, U. Tennessee, Knoxville.

88 PPM HCl/pH₂



Gas Phase (HCl)₂

High resolution, jet-cooled infrared spectroscopy of (HCl)₂: Analysis of ν_1 and ν_2 HCl stretching fundamentals, interconversion tunneling, and mode-specific predissociation lifetimes

Michael D. Schuder,^{a)} Christopher M. Lovejoy,^{b)} Robert Lascola,^{c)} and David J. Nesbitt^(d)
*Joint Institute for Laboratory Astrophysics, National Institute of Standards and Technology and
University of Colorado, and the Department of Chemistry and Biochemistry, University of Colorado,
Boulder, Colorado 80309*

(Received 5 April 1993; accepted 7 June 1993)

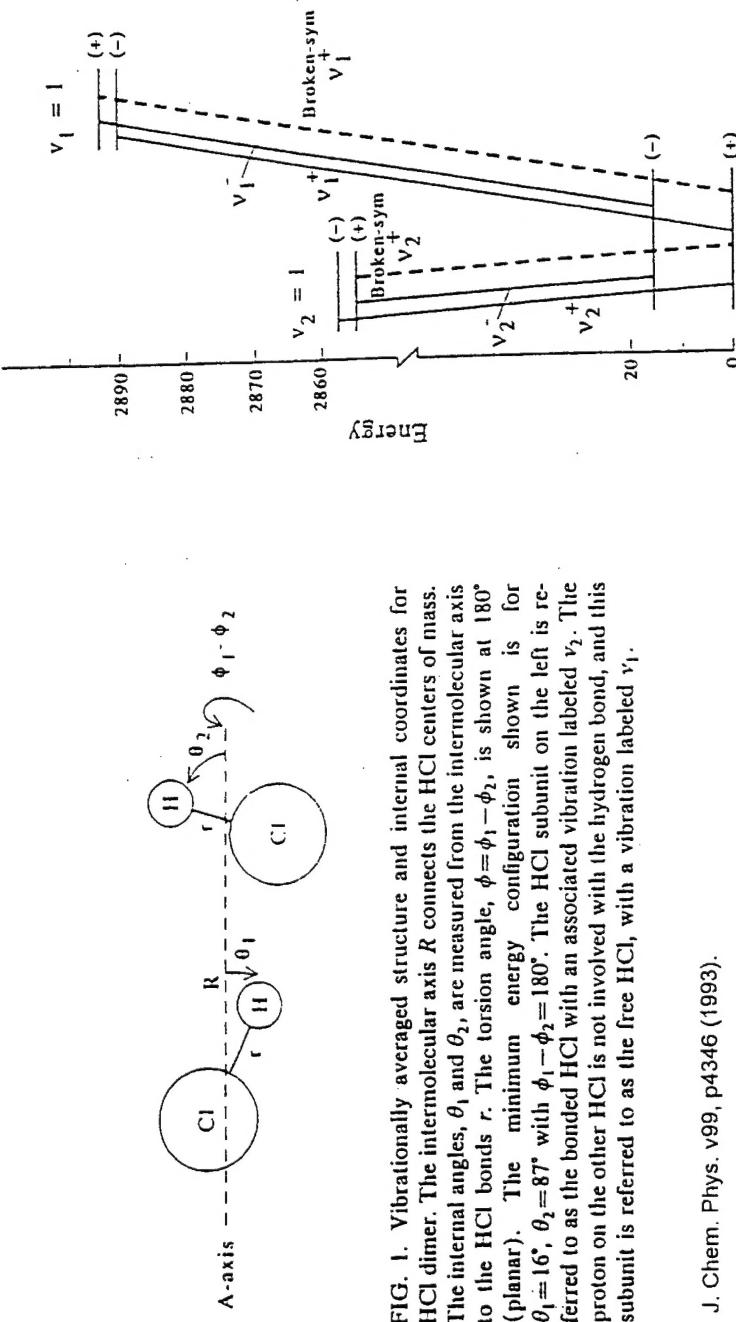
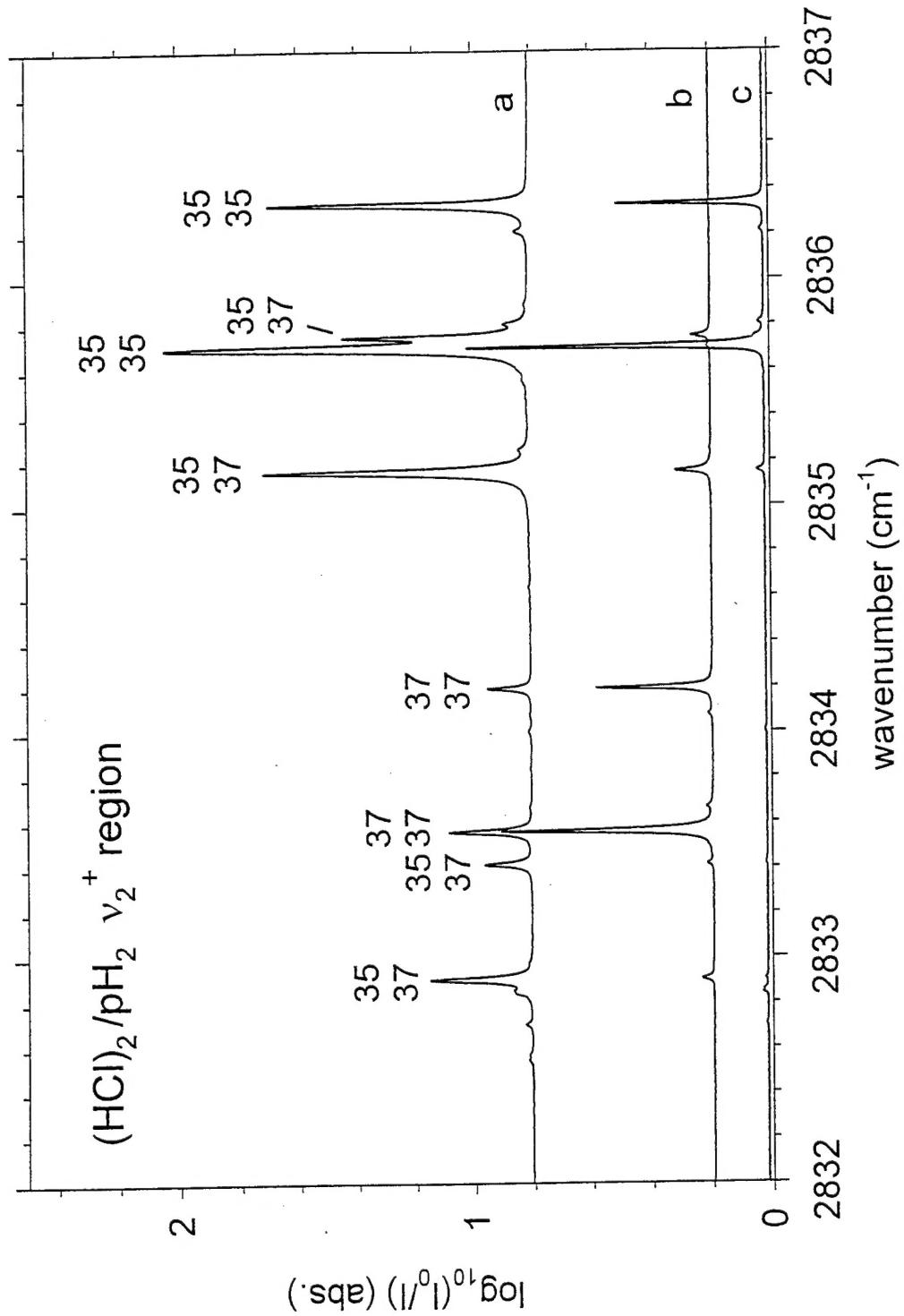


FIG. 1. vibrationally averaged structure and internal coordinates for HCl dimer. The intermolecular axis R connects the HCl centers of mass. The internal angles, θ_1 and θ_2 , are measured from the intermolecular axis to the HCl bonds r . The torsion angle, $\phi = \phi_1 - \phi_2$, is shown at 180° (planar). The minimum energy configuration shown is for $\theta_1 = 16^\circ$, $\theta_2 = 87^\circ$ with $\phi_1 - \phi_2 = 180^\circ$. The HCl subunit on the left is referred to as the bonded HCl with an associated vibration labeled ν_2 . The proton on the other HCl is not involved with the hydrogen bond, and this subunit is referred to as the free HCl, with a vibration labeled ν_1 .

(HCl)₂/pH₂ isotopomers

Department of Chemistry, University of Wyoming, Laramie, WY, 20 April 2001



analysis in collaboration with D.T. Anderson, U. Wyoming.

HEDM Cryosolids Accomplishments

(a list of “things that’ll never work.”)

- * Trapped Li, B, Na, Mg, Al atoms in solid hydrogen at $T \approx 2$ K; attempts to demonstrate useful chemical energy storage still in progress
- * Demonstrated production of gram-scale optically transparent pH₂ solids by rapid vapor deposition.
- * Demonstrated that vapor deposited pH₂ solids are densest close-packed solids, NOT amorphous.
- * Demonstrated suitability of vapor deposited pH₂ solids as hosts for high resolution IR absorption spectroscopy of chemically interesting dopants; spectral assignments ongoing.
- * Generalized phenomena of dopant-induced and co-operative IR absorptions to chemically interesting dopants.